



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

re the Application of:

Jun Feng et al.

Serial No.: 10/809,635

Filed: March 24, 2004

For: DIPEPTIDYL PEPTIDASE
INHIBITORS

Group Art Unit: 1646

Examiner: Not Yet Assigned

INFORMATION DISCLOSURE STATEMENT

Mail Stop Amendment
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

In accordance with 37 CFR §§ 1.97 and 1.98, the items identified in this Information Disclosure Statement ("IDS") are brought to the attention of the Office. The items are listed on the attached form PTO-1449, and copies are included for the Examiner's convenience. As the Office no longer requires copies of U.S. patents and applications, the U.S. copies are not being submitted with this IDS. However, if the Examiner would like copies of the U.S. cited references, Applicants will provide the references upon request. The item listed on the attached form PTO-1449 at "EN" is a non-English language article. In accordance with 37 CFR § 1.98(a)(3)(i), the following is a concise explanation of the relevance of this article:

The article by P.O. Bezuglyi relates to the synthesis of arylsulfonfyl hydrazides of 3-R-quinazalone-4-carbonyl-2-acid.

The items identified in this IDS may or may not be "material" pursuant to 37 CFR § 1.56. The submission thereof by Applicant is not to be construed as an admission that any such patent, publication or other information referred to therein is material or considered to be material (37 CFR § 1.97(h)), or even qualifies as "prior art" under 35 USC § 102 with respect to this invention unless specifically designated by Applicants as such.

INFORMATION DISCLOSURE STATEMENT FILING PROVISION:

☒ This IDS is believed to be timely in that it is being submitted under 37 CFR § 1.97(b), that is (1) within three months of the filing date of the application, which is not a continued prosecution application filed under § 1.53(d); or (2) within three months of entry of the national stage as set forth in 37 CFR § 1.491; or (3) before the mailing of a first Office action on the merits; or (4) before the mailing of a first Office action after filing a request for continued examination under § 1.114. Thus, no fee is required.

☒ However, if the undersigned is in error in this regard, Applicant respectfully requests that the Office consider this IDS as filed under 37 CFR § 1.97(c), if applicable, and charge the fee due under 37 CFR § 1.17(p) to the deposit account referenced below.

☐ However, if the undersigned is in error in this regard, Applicant respectfully requests that the Office consider this IDS as filed under 37 CFR § 1.97(c), if applicable, and a statement under 37 CFR § 1.97(e) is included below, thus no fee is required.

☐ This IDS is being submitted under 37 CFR § 1.97(c), that is after mailing of a first Office action on the merits, but before a Final Action under 37 CFR § 1.113 or a Notice of Allowance under 37 CFR § 1.311.

☐ The fee due under 37 CFR § 1.17(p) is submitted herewith.

☐ A statement under 37 CFR § 1.97(e) is included below, thus no fee is required. In the event that this IDS is not received before a Final Action or a Notice of Allowance, then Applicant respectfully requests that the Office consider the filing of these papers to be submitted under 37 CFR § 1.97(d) and charge the fee due under 37 CFR § 1.17(p) to the deposit account below.

☐ This IDS is being submitted under 37 CFR § 1.97(d), that is after a Final Action under 37 CFR § 1.113 or a Notice of Allowance under 37 CFR § 1.311, but before payment of the issue fee. A statement under 37 CFR § 1.97(e) is included below. The fee due under 37 CFR § 1.17(p) is submitted herewith.

STATEMENT UNDER 37 CFR § 1.97(e):

☐ Each item contained in this IDS was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of this IDS.

☐ No item contained in this IDS was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing

Patent
SYR-DPP-IV-5004-U

this statement after making reasonable inquiry, no item of information contained in this IDS was known to any individual designated in 37 CFR § 1.56(c) more than three months prior to the filing of this IDS.

PAYMENT AND/OR AUTHORIZATION TO CHARGE FEES:

- ☐ A check in the amount of _____ is enclosed for the above fee(s).
☐ Please charge to Deposit Account No. **50-2256** for the above fee(s).

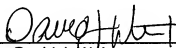
Although Applicants do not believe any fees are required, the Commissioner is authorized to charge any fees required by the filing of these papers to Syrrx's Deposit Account No. **50-2256**.

Respectfully submitted,

SYRRX, INC.

Dated: February 18, 2005

By: _____


David J. Weitz
Reg. No. 38,362

Customer No. **32793**
Syrrx, Inc.
10410 Science Center Drive
San Diego, CA 92121
Tel: (858) 622-8528
Fax: (858) 550-0992

PTO/SB/08A (08-03)

Approved for use through 07/31/2008. OMB 0651-0031

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

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Substitute for form 1449/APTO

**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

(use as many sheets as necessary)

Sheet 1 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

**U.S. PATENT DOCUMENTS**

Examiner Initials *	Cite No. 1	Document Number	Publication Date/ Issue Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number - Kind Code ² (if known)			
/KH/	AA	US1974/3623135	07-09-1974	Pilgram et al.	
	AB	US1996/5512549	04-30-1996	Chen et al.	
	AC	US1996/5580979	12-03-1996	Bachovchin	
	AD	US1997/5614492	03-25-1997	Habener	
	AE	US2000/6156739	12-5-2000	Griffin et al.	
	AF	US2000/6166063	12-26-2000	Villhauer	
	AG	US2001/6258597-B1	07-10-2001	Bachovchin	
	AH	US2001/0020008-A1	09-06-2001	Demuth et al.	
	AI	US2001/6303661-B1	10-16-2001	Demuth et al.	
	AJ	US2001/6319893-B1	11-20-2001	Demuth et al.	
	AK	US2001/0051646-A1	12-13-2001	Demuth et al.	
	AL	US2002/0049153-A1	04-25-2002	Bridon et al.	
	AM	US2002/0049164-A1	04-25-2002	Demuth et al.	
	AN	US2002/6380398-B2	04-30-2002	Kanstrup et al.	
	AO	US2002/0082427-A1	06-27-2002	Demuth et al.	
	AP	US2002/6448045-B1	09-10-2002	Levine et al.	
	AQ	US2002/0198242-A1	12-26-2002	Demuth et al.	
	AR	US2002/0198380-A1	12-26-2002	Belzer et al.	
	AS	US2002/6500804-B2	12-31-2002	Demuth et al.	
	AT	US2003/0008925-A1	01-09-2003	Demuth et al.	
	AU	US2003/6548481-B1	04-15-2003	Demuth et al.	
V	AV	US2003/0092630-A2	05-15-2003	Demuth et al.	
	AW	US2003/0119750-A1	06-26-2003	Demuth et al.	
	AX	US2003/0130199-A1	07-10-2003	von Hoersten et al.	
	AY	US2003/0134802-A1	07-17-2003	Demuth et al.	
	AZ	US2003/0135023-A1	07-17-2003	Demuth et al.	
	BA	US2003/0148961-A1	08-07-2003	Heiser et al.	
	BB	US2003/0153509-A1	08-14-2003	Bachovchin et al.	
	BC	US2003/0162820-A1	08-28-2003	Demuth et al.	
	BD	US2003/0166578-A1	09-04-2003	Arch et al.	
	BE	US2003/6620910-B1	09-16-2003	Calas et al.	
V	BF	US2003/0176357-A1	09-18-2003	Pospisilik et al.	
	BG	US2003/0199451-A1	10-23-2003	Mogensen et al.	
	BH	US2003/0199672-A1	10-23-2003	Knudsen et al.	

Examiner Signature	/Kahsay Habte/	Date Considered	08/14/2008
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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 Applicant's unique citation designation number (optional). 2 Applicant is to place a check mark here if English language Translation is attached. This collection of information is required by 37 CFR 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

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Sheet 2 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	BI	US2003/0236272-A1	12-25-2003	Richard David Carr	
	BJ	US2004/6703238-B2	03-09-2004	Bachovchin	
	BK	US2004/0054171-A1	03-18-2004	Jensen et al.	
	BL	US2004/0058876-A1	03-25-2004	Hoffmann et al.	
	BM	US2004/0132732-A1	07-08-2004	Han et al.	
	BN	US2004/0167191-A1	06-26-2004	Demuth et al.	
	BO	US2004/0171555-A1	09-02-2004	Demuth et al.	

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Foreign Patent Document Country Code ² - Number ³ - Kind Code ⁴ (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁵
/KH/	BP	FR 2,162,106 (English Abstract-1973)	11-30-1972	Amschler et al.		
	BQ	WO 89/10701	11-16-1989	BASF		
	BR	EP 0378255-A2	07-18-1990	Janssen Pharmaceutica		
	BS	GB 2230527-A	10-24-1990	Imperial Chemical Industries Plc		
	BT	WO 91/12001	08-22-1991	Merck & Co., Inc.		
	BU	WO 93/21162	01-28-1993	Nissan Chemical Industries, Ltd.		
	BV	WO 93/08259 (A2)	04-29-1993	New England Medical Center Hospitals, Inc.		
	BW	WO 93/08259 (A3)	04-29-1993	New England Medical Center Hospitals, Inc.		
	BX	EP 0547442-A1	06-23-1993	E.R. Squibb & Sons, Inc.		
	BY	WO 94/03055	02-17-1994	U.S. Government, Secty. HHS		
	BZ	EP 0587377-A2	03-16-1994	Eli Lilly and Company		
	CA	WO 95/35031	12-28-1995	La Trobe University		
	CB	WO 96/32384	10-17-1996	Talho Pharmaceutical Co., Ltd.		
	CC	WO 96/38550	12-05-1996	Dana-Farber Cancer Institute, Inc.		
	CD	WO 97/40832	11-06-1997	Hans-Knoll-Institut Fur Naturstoff		
	CE	JP 9295977	11-18-1997	Tenumo Corp.		
	CF	WO 98/00439	01-08-1998	Trustees of Tufts College		
	CG	WO 98/24780	06-11-1998	Amgen Inc.		
	CH	WO 99/16864	04-08-1999	Point Therapeutics, Inc.		
	CI	WO 99/38501	06-05-1999	Trustees of Tufts University		

Examiner Signature	/Kahsay Habte/	Date Considered	08/14/2008
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**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

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Sheet 3 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	CJ	WO 99/50249	10-07-1999	Du Pont Pharmaceuticals Company	
	CK	WO 99-61431	12-02-1999	Probiolog	
	CL	WO 99/67278	12-29-1999	Pro-Biolog	
	CM	WO 99/67279	12-29-1999	Pro-Biolog	
	CN	WO 00/07617	02-17-2000	Novo Nordisk	
	CO	WO 00/09666	02-24-2000	U.S. Government, Secty. HHS	
	CP	WO 00/15211	03-23-2000	Akasis Pharmaceuticals, Inc.	
	CQ	WO 00/76986-A1	04-11-2000	Probiolog	
	CR	WO 00/34241	06-15-2000	Novartis AG	
	CS	WO 00/47219	08-17-2000	Ontogeny, Inc.	
	CT	WO 00/53171	09-14-00	Moteni L. E. C. Dei Fratelli Alitti Societa' Di Esercizio S.P.A.	
	CJ	WO 00/57721	10-05-2000	Akasis Pharmaceuticals, Inc.	
	CV	WO 01/14318-A2	03-01-2001	Probiolog	
	CW	WO 01/34594-A1	05-17-2001	Gulford Pharmaceuticals, Inc.	
	CX	WO 01/52825-A2	07-26-2001	Novartis AG	
	CY	WO 01/56988-A1	08-09-2001	Kirin Beer Kabushiki Kaisha	
	CZ	WO 01/70729-A1	09-27-2001	Sanofi-Synthelabo	
	DA	WO 01/97808-A1	12-27-2001	Smithkline Beecham PLC	
	DB	WO 02/34242-A2	05-02-2002	Probiolog AG	
	DC	WO 02/34243-A2	05-02-2002	Probiolog AG	
	DD	WO 02/083109-A1	10-24-2002	Ferring BV	
	DE	JP 2002/338466	11-27-2002	Tanabe Seiyaku Co Ltd	
	DF	WO 03/002593-A2	01-09-2003	Probiolog AG	
	DG	WO 03/002595-A2	01-09-2003	Probiolog AG	
	DH	WO 03/002596-A2	01-09-2003	Probiolog AG	
	DI	WO 03/016335-A2	02-27-2003	Probiolog AG	
	DJ	WO 03/022871-A2	03-20-2003	Probiolog AG	
	DK	WO 03/026652-A1	04-03-2003	Bristol-Myers Squibb Company	
	DL	WO 03/030946-A1	04-17-2003	Novartis AG	
	DM	WO 03/033524-A2	04-24-2003	Probiolog AG	
	DN	JP 2003/128551	05-08-2003	Sankyo Co LTD	
	DO	WO 03/040174-A2	05-15-2003	Probiolog AG	
	DP	WO 03/045228-A2	06-05-2003	Trustees of Tufts College	
	DQ	WO 03/045977-A2	06-05-2003	Trustees of Tufts College	
	DR	WO 03/048081-A2	06-12-2003	Bristol-Myers Squibb Company	
	DS	WO 03/048158-A1	06-12-2003	Bristol-Myers Squibb Company	
	DT	WO 03/057200-A2	07-17-2003	Novo Nordisk	

Examiner
Signature

/Kahsay Habte/

Date

Considered

08/14/2008

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U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 4 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	DU	WO 03/063903-A2	06-07-2003	Probioludng AG	
	DV	WO 03/072556-A1	09-04-2003	Probioludng AG	
	DW	WO 03/082898-A2	10-09-2003	Probioludng AG	
	DX	WO 03/092605-A2	11-13-2003	Trustees of Tufts College	
	DY	WO 03/099279-A1	12-04-2003	Novartis AG	
	DZ	WO 03/098818-A1	12-04-2003	Chiron Corporation	
	EA	WO 03/106416-A2	12-24-2003	Smithkline Beecham Corporation	
	EB	WO 2004/017989-A1	03-04-2004	Probioludng AG	
	EC	JP 2004/09600-A	04-02-2004	Sankyo Co. Ltd.	
	ED	WO 2004/031374-A2	04-15-2004	Probioludng AG	
	EE	JP 2004/123738-A	04-22-2004	Takeda Chem Ind Ltd	
	EF	WO 2004/037176-A2	05-06-2004	Bristol-Myers Squibb Company	

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T 2
/KH/	EG	ARGAUD, DORIANE et al., Metformin decreases gluconeogenesis by enhancing the pyruvate kinase flux in isolated rat hepatocytes, European J. Biochem. 213, 1341-1348 (1993).	
	EH	ASHCROFT, STEPHEN J.H. et al., Structure-activity relationships of alloxan-like compounds derived from uric acid, Br. J. Pharmac. (1986), 89 pp. 469-472.	
	EI	BAL, GUNTHER, Dipeptidyl Peptidase IV and Prolyl Oligopeptidase: Design, Synthesis and Evaluation of Substrates and Inhibitors, (2002) Universiteit Antwerpen.	
	EJ	BARAKAT, S.E.S., Synthesis and hypoglycemic activity of some new 3-(4-(((cydohexylamino) carbonyl) amino)sulfonylphenyl)-(4(3H)-quinazolinones, Az. J. Pharm. Sci., Vol. 25, (2000), pp. 48-57.	
	EK	BARAKAT, S.E.S., Synthesis and Hypoglycemic Activity of Some New 4(3H)-Quinazolinone Analogues, Saudi Pharmaceutical Journal, Vol. 8, No. 4 (2000) pp. 198-204.	
	EL	BAKER, B.R. et al., Irreversible Enzyme Inhibitors. On the Mode of Pyrimidine Binding of 5-alkyl and 5-Arylpyrimidines to Dihydrofolate Reductase (1,2), Journal of Heterocyclic Chemistry Vol. 4 (1967) pp. 39-48.	
	EM	BELGODERE, ELENA et al., Synthesis of Substituted Pyrimidines, Study of the Structure and of the Tautomeric Equilibria, (1976) Chem. Abstracts, Columbus, OH Vol. 85 No. 9.	
	EN	BEZUGLYI, P.O. et al., Synthesis of arylsulfonyl hydrazide of 3-R-quinazolinone-4-carboxyl-2-acid, Pharmaceutical Journal (1979), pp. 70-71.	

Examiner
Signature

/Kahsay Habte/

Date
Considered

08/14/2008

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

(use as many sheets as necessary)

Sheet 5 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	EO	BHADURI, A.P. et al., Urinary Metabolite of 2-Piperazino-3 (H)-4-Quinazolinone (Centiperalone), A Potent Blood Sugar Lowering Agent, Indian J. Biochem. Biophys., Vol. 12 (1975), pp. 413-414.
	EP	BOURAS, MOHAMMED, et al., Metabolism of enterostatin in rat intestine, brain, membranes and serum: differential involvement of proline-specific peptidases, Peptides, Vol. 16, No. 3, (1995), pp. 399-405.
	EQ	BRUN, JEAN-FREDERIC, et al., Effects of Oral Zinc Gluconate on Glucose Effectiveness and Insulin Sensitivity in Humans, Biological Trace Element Research Vol. 47 (1995), pp. 385-391.
	ER	BUCKLEY, DI, Analysis of the Degradation of Insulintropin [GLP-1 (7-37)] in Human Plasma and Production of Degradation Resistant Analogs.
	ES	CHATTERJEE, A.K. et al., Effect of Centiperalone in Insulin Deficient Diabetes, Indian Journal of Experimental Biology Vol. 18 (1980), pp. 1005-1006.
	ET	CHATTERJEE, A.K. et al., Effect of Centiperalone, a New Hypoglycemic Agent on Insulin Biosynthesis & Release from Isolated Pancreatic Islets of Rat, Indian Journal of Experimental Biology Vol. 20 (1981) pp.270-272.
	EU	COPPOLA, GARY M. et al., 1-Aminomethylisoquinoline-4-carboxylates as Novel Dipeptidylpeptidase IV Inhibitors, Bioorganic & Medicinal Chemistry Letters Vol. 10 (2000), pp. 1555-1558.
	EV	DEACON, CAROLYN F. et al., Degradation of Glucagon-Like Peptide 1 in Vitro Yields an N-Terminally Truncated Peptide That is a Major Endogenous Metabolite in Vivo, Journal of Clinical Endocrinology and Metabolism Vol. 80, No. 3 (1995), pp. 952-957.
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✓	FC	FANTUS, I. GEORGE, et al., Mechanism of Action of Metformin: Insulin Receptor and Postreceptor Effects in Vitro and in Vivo, J. Clinical Endocrinology & Metabolism (1986), pp. 898-905.

Examiner Signature	/Kahsay Habte/	Date Considered	08/14/2008
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Sheet 6 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	FD	GARRATT, PETER J. et al., A Novel Synthesis of Dihydropyrimidines, J. Chem. Soc., Chem. Commun. (1987), pp.568-569.
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Examiner Signature	/Kahsay Habte/	Date Considered	08/14/2008
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Sheet 7 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	FS	KUSAR, MIHAEL et al., Diethyl N,N-Dimethylaminomethylmaleonate in the Synthesis of Fused Heterocyclic Systems, <i>Heterocyclic Chem.</i> 33 (1996) pp. 1041-1046.	
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Examiner Signature	/Kahsay Habte/	Date Considered	08/14/2008
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Sheet 8 of 10

Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	GH	MUKHERJEE, SURATH K. et al., Influence of Timing Oral Dosing of a Novel Hypoglycaemic Agent A-4166 in Relation to Food, <i>Diabetologia</i> Vol. 38 A194 Supplement 1 (1995).	
	GI	MUKHERJEE, Subal S. et al., Studies on the Mechanism of Centipalalone-Induced Hypoglycemia, <i>Acta Diabet.</i> Lat. 13, 8, (1976) pp. 8-19.	
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	GS	POJE, M. et al., Diabetogenic action of alloxan-like derivatives of uric acid, <i>Experientia</i> 36 (1980) pp. 78-9.	
	GT	POJE, M. et al., Oxidation of Uric Acid. 4. Synthesis, Structure, and Diabetogenic Action of 5-Imino-2,4,6 (1H,3H,5H)-pyrimidinetrione Salts and Their Alloxan-like Covalent Adducts, <i>J. Med. Chem.</i> 26 (1983) 861-4.	
	GU	POLACEK, I. et al., Hypoglycemic Activity of Amine Derivatives, <i>Arzneim.-Forsch./Drug Res.</i> 26 (1976), 791-93.	
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**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

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Complete if Known

Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

Sheet 9 of 10

/KH/	GW	RAM, VISHNU JI et al., Synthesis and Antihyperglycemic Activity of Suitably Functionalized 3H-quinoxalin-4-ones, Bioorganic & Medicinal Chemistry 11 (2003), pp. 2430-2444.
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	HC	SETH, M. et al., Syntheses of 2-Substituted & 2,3-Distributed 4(3H)-Quinoxalones, Indian Journal of Chemistry, Vol. 14B (1975), 536-540.
	HD	SHIMAZAWA, RUMIKO et al., Novel Small Molecule Nonpeptide Aminopeptidase N Inhibitors with A Cyclic Imide Skeleton, J. Enzyme Inhibition, Vol. 14 (1999) pp. 259-275.
	HE	SHISHEVA, ASSIA et al., Insulinlike Effects of Zinc Ion in Vitro and in Vivo; Preferential Effects on Desensitized Adipocytes and Induction of Normoglycemia in Streptozotocin-Induced Rats, Diabetes, Vol. 41 (1992), pp. 982-988.
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↓	HK	VILLHAUER, EDWIN B. et al., 1-[[[3-Hydroxy-1-adamantyl]amino]acetyl]-2-cyano-(5-pyrrolidine): A Potent, Selective, and Orally Bioavailable Dipeptidyl Peptidase IV Inhibitor with Antihyperglycemic Properties, J. Med. Chem. 46 (2003), pp. 2774-2789.

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Sheet 10 of 10

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Application Number	10/809,635
Filing Date	March 24, 2004
First Named Inventor	Jun Feng
Group Art Unit	1646
Examiner Name	Not Yet Assigned
Attorney Docket Number	SYR-DPP-IV-5004-U

/KH/	HL	WELLS, CAROL L. et al., Role of Anaerobic Flora in the Translocation of Aerobic and Facultatively Anaerobic Intestinal Bacteria, Infection and Immunity, Vol. 55, No. 11 (1987) pp. 2689-94.	
	HM	WIEDEMAN, PAUL E. et al., Dipeptidyl peptidase IV inhibitors for the treatment of impaired glucose tolerance and type 2 diabetes, Current Opinion in Investigational Drugs, Vol. 4, No. 4 (2003), pp. 412-420.	
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✓	HQ	ZHANG, ANQI et al., Vanadate Stimulation of Insulin Release in Normal Mouse Islets, The Journal of Biological Chemistry, Vol. 266, No. 32 (1991), pp. 21649-56.	

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1 Applicant's unique citation designation number (optional). 2 Applicant is to place a check mark here if English language Translation is attached. This collection of information is required by 37 CFR 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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